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AMENDMENTS TO THE CLAIMS

Please amend the claims to read as follows:

- 1. (Amended) A method of treating a patient, comprising the step of delivering to [the] <u>said</u> patient's cardiac autonomic structures <u>a</u> nucleic acid <u>molecule</u> which, when expressed, increases nitric oxide synthase levels.
- 2. (Amended) A pharmaceutical composition comprising a The use of nucleic acid molecule which, when expressed in cardiac autonomic structures, increases nitric oxide synthase levels, in the manufacture of a medicament.
- 3. (Amended) The method or use of any preceding of claim 1, wherein treatment is for increasing cardiac vagal tone [and/or], increasing cardiac vagal responsiveness, [for] increasing bradycardia, [for] reducing cardiac autonomic impairment, [for] reducing the risk of sudden cardiac death, [for] reducing arrhythmia (e. g. atrial fibrillation and/or ventricular arrhythmia), [for] reducing the risk of myocardial infarction[, and/] or [for] reducing hypertension.
- 4. (Amended) The method [or use] of any preceding claim 1, wherein said nucleic acid molecule is delivered to the vagus nerve.
- 5. (Amended) The method [or use] of any preceding claim 1, wherein [the] said nitric oxide synthase is NOS-1 [and/]or NOS-3.
- 6. (Amended) The method [or use] of claim 5, wherein the NOS-1 is human NOS-1.
- 7. (Amended) The method [or use] of any preceding claim 1, wherein [the] said nucleic acid molecule is targeted to cardiac tissue.
- 8. (Amended) The method [or use] of any preceding claim 1, wherein expression of the nitric oxide synthase from [the] said nucleic acid molecule is regulated by a non-constitutive promoter.

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- 9. (Amended) The method [or use] of any one of claims claim 1 [to 7], wherein expression of [the] said nitric oxide synthase is regulated by a promoter which is specifically active in cholinergic ganglia tissue.
- 10. (Amended) The method [or use] of any-preceding claim 1, wherein [the] said nucleic acid molecule is packaged within a non-viral gene therapy vector.
- 11. (Amended) The method [or use] of claim 10, wherein [the] <u>said</u> nucleic acid <u>molecule</u> is delivered as naked DNA.
- 12. (Amended) The method [or use] of any preceding claim 3, wherein cardiac vagal tone in the patient is increased by at least 10%.
- 13. (Amended) The method [or use] of any preceding claim 1, wherein [the] said nucleic acid molecule comprises DNA.
- 14. (Amended) The method [or use] of any preceding claim 1, wherein [the] said nucleic acid molecule is non-replicating.
- 15. (Amended) The method [or use] of any preceding claim 1, wherein [the] said nucleic acid molecule is non-integrating.
- 16. (Amended) The method [or use] of any preceding claim 1, wherein [the] said nucleic acid molecule is an autonomously replicating episomal or extrachromosomal vector, such as a plasmid.
- 17. (Amended) The method [or use] of any preceding claim 1, wherein [the] said nucleic acid molecule is delivered to the heart within microbubbles which can be disrupted by ultrasound.
- 18. (Amended) A nucleic acid molecule comprising a non-constitutive promoter and a coding sequence, wherein: (a) [the] <u>said</u> promoter is operably linked to the coding sequence to control transcription of the coding sequence; (b) [the] <u>said</u> promoter is specifically active in cholinergic ganglia tissue; and (c) [the] <u>said</u> coding sequence encodes a nitric oxide synthase.

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19. (Amended) A pharmaceutical composition comprising the nucleic acid <u>molecule</u> of claim 18, for use as a medicament.